

chain nodes :

22

ring nodes :

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21

chain bonds :

6-16 11-15

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12 13-14 13-17 13-21  
14-15 15-16 16-17 17-18 18-19 19-20 20-21

exact/norm bonds :

13-14 13-17 13-21 14-15 17-18 18-19 19-20 20-21

exact bonds :

6-16 11-15 15-16 16-17

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 7-8 7-12 8-9 9-10 10-11 11-12

isolated ring systems :

containing 1 : 7 : 13 :

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom 11:Atom  
12:Atom 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom 19:Atom 20:Atom 21:Atom  
22:CLASS 23:CLASS

=> d his

(FILE 'HOME' ENTERED AT 15:07:24 ON 20 NOV 2001)

FILE 'REGISTRY' ENTERED AT 15:07:30 ON 20 NOV 2001  
L1                   STRUCTURE UPLOADED  
L2                   8 S L1  
L3                   160 S L1 FULL

FILE 'CA' ENTERED AT 15:08:27 ON 20 NOV 2001  
L4                   5 S L3  
L5                   0 S L4 AND BAXTER, I?/AU  
L6                   0 S L4 AND PD < DECEMBER 1998

FILE 'CAOLD' ENTERED AT 15:10:02 ON 20 NOV 2001

=> s 13

L7                   0 L3

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PASSWORD:  
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=> file eg

'EG' IS NOT A VALID FILE NAME  
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=> file req

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.15	0.15

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STRUCTURE FILE UPDATES: 19 NOV 2001 HIGHEST RN 371110-14-4  
DICTIONARY FILE UPDATES: 19 NOV 2001 HIGHEST RN 371110-14-4

TSCA INFORMATION NOW CURRENT THROUGH July 7, 2001

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:  
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

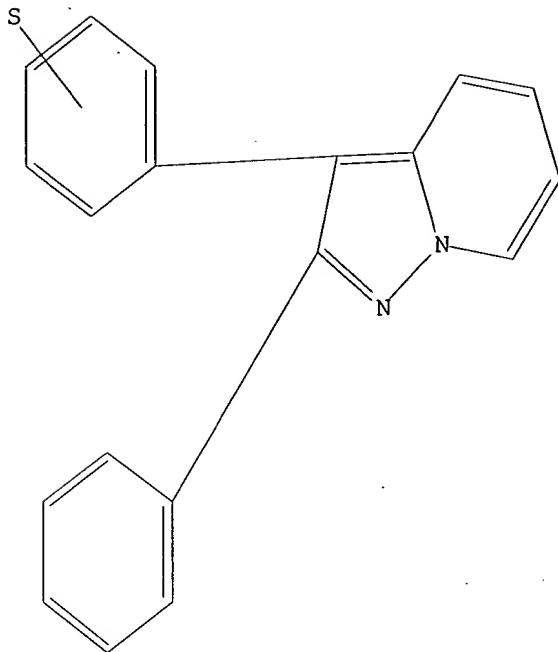
⇒

Uploading 09830836.str

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS  
L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 15:08:14 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED - 53 TO ITERATE

100.0% PROCESSED 53 ITERATIONS 8 ANSWERS  
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE \*\*COMPLETE\*\*  
BATCH \*\*COMPLETE\*\*  
PROJECTED ITERATIONS: 624 TO 1496  
PROJECTED ANSWERS: 8 TO 329

L2 8 SEA SSS SAM L1

=> s 11 full

THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 133.25 U.S. DOLLARS  
DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y

FULL SEARCH INITIATED 15:08:21 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED - 1167 TO ITERATE

100.0% PROCESSED 1167 ITERATIONS 160 ANSWERS  
SEARCH TIME: 00.00.02

L3 160 SEA SSS FUL L1

=> file.ca

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	133.87	134.02

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FILE COVERS 1947 - 15 Nov 2001 VOL 135 ISS 22  
 FILE LAST UPDATED: 15 Nov 2001 (20011115/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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=> s 13

L4 5 L3

=> s 14 and baxter, i?/au

L5 86 BAXTER, I?/AU  
 0 L4 AND BAXTER, I?/AU

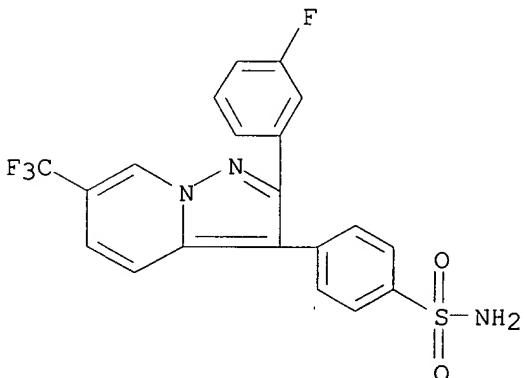
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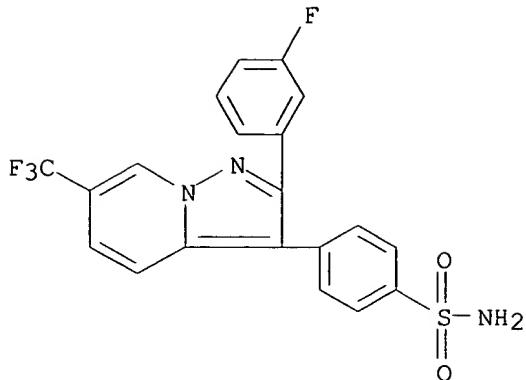
L6 17147450 PD < DECEMBER 1998  
 (PD<19981200)  
 0 L4 AND PD < DECEMBER 1998

=> d 14, ibib abs fhitstr, 1-5

L4 ANSWER 1 OF 5 CA COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 135:147440 CA  
 TITLE: Use of cyclooxygenase-2 (COX-2) inhibitors as  
 gastropokinetic agents  
 INVENTOR(S): Mangel, Allen Wayne; Naylor, Alan  
 PATENT ASSIGNEE(S): Glaxo-Smithkline, UK  
 SOURCE: PCT Int. Appl., 28 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001056573	A1	20010809	WO 2001-GB423	20010201
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
PRIORITY APPLN. INFO.:			GB 2000-2336	A 20000201
AB	The invention provides a COX-2 inhibitor or a pharmaceutically acceptable deriv. thereof for use in the treatment of a disorder ameliorated by a gastropokinetic agent.			
IT	<b>267235-56-3</b>			
	RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (cyclooxygenase-2 inhibitors as gastropokinetic agents)			
RN	267235-56-3 CA			
CN	Benzenesulfonamide,			
	4-[2-(3-fluorophenyl)-6-(trifluoromethyl)pyrazolo[1,5-a]pyridin-3-yl]- (9CI) (CA INDEX NAME)			





REFERENCE COUNT:

6

REFERENCE(S):

P47

(1) Hayakawa, T; J SMOOTH MUSCLE RES 1999, V35(2),

MEDLINE  
 (3) Merck Sharp & Dohme; GB 2325161 A 1998 CA  
 (4) Morgan, G; EUROPEAN JOURNAL OF GASTROENTEROLOGY  
 AND HEPATOLOGY 1999, V11(4), P393 CA  
 (5) Panacea Biotec Ltd; EP 0812591 A 1997 CA  
 (6) Pozen Inc; WO 0048583 A 2000 CA  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 5 CA COPYRIGHT 2001 ACS

ACCESSION NUMBER: 135:147439 CA

TITLE: Use of cyclooxygenase-2 (COX-2) inhibitors for constipation

INVENTOR(S): Mangel, Allen Wayne; Naylor, Alan

PATENT ASSIGNEE(S): Glaxo Group Limited, UK

SOURCE: PCT Int. Appl., 21 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

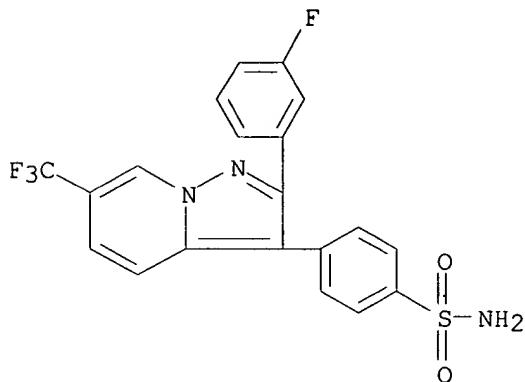
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001056555	A2	20010809	WO 2001-GB416	20010201
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.: GB 2000-2312 A 20000201

AB The invention provides a COX-2 inhibitor or a pharmaceutically acceptable deriv. thereof for use in the treatment of constipation.

IT 267235-56-3

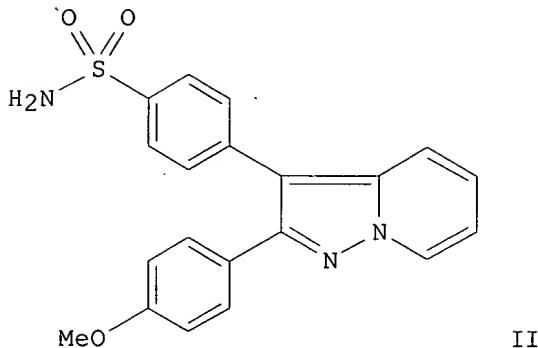
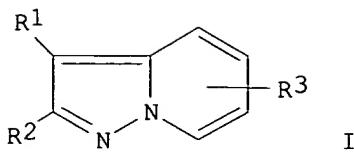
RL: BAC (Biological activity or effector, except adverse); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
 (cyclooxygenase-2 inhibitors for treatment of constipation)  
 RN 267235-56-3 CA  
 CN Benzenesulfonamide,  
 4-[2-(3-fluorophenyl)-6-(trifluoromethyl)pyrazolo[1,5-a]pyridin-3-yl]- (9CI) (CA INDEX NAME)



L4 ANSWER 3 OF 5 CA COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 135:5612 CA  
 TITLE: Preparation of new pyrazolo terpyridines as remedies  
 for inflammation, autoimmune diseases  
 INVENTOR(S): Yamamoto, Hiroyuki; Takahashi, Fumie; Kato, Takeshi;  
 Nakamura, Katsuya; Manabe, Koji  
 PATENT ASSIGNEE(S): Fujisawa Pharmaceutical Co., Ltd., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 64 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2001139575	A2	20010522	JP 1999-323692	19991115

GI



AB The pyrazolo terpyridine or that salt which is cyclooxygenase - 2 (COX-II) inhibitors, those prodn. methods, the medicine compn., and the person or the animal which contain those inflammation condition, u painfully, prevention of the autoimmune disease and / or the method of treating is offered. Below-mentioned general formula (I) [ in the formula, the R1 and

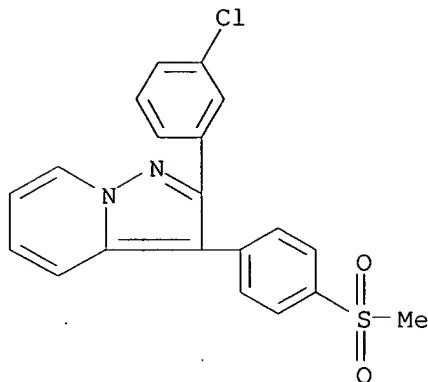
the R2, the resp. hydrogen, the hydrogen, the low-grade alkyl group and the halogen et cetera, mean, R3 such as low-grade alkyl group and the cyclo (low grade) alkyl group resp. ] So the chem. compd. which is displayed or that salt.

IT 340321-35-9P

RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
(prepn. of new pyrazolo terpyridines as remedies for inflammation autoimmune diseases)

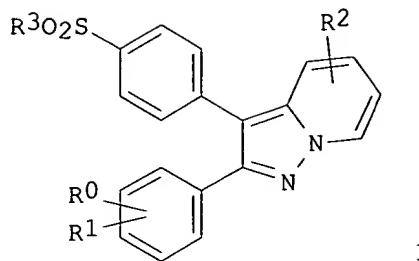
RN 340321-35-9 CA

CN Pyrazolo[1,5-a]pyridine, 2-(3-chlorophenyl)-3-[4-(methylsulfonyl)phenyl]- (9CI) (CA INDEX NAME)



L4 ANSWER 4 OF 5 CA COPYRIGHT 2001 ACS  
 ACCESSION NUMBER: 133:222726 CA  
 TITLE: Preparation of pyrazolopyridines as selective inhibitors of COX-2  
 INVENTOR(S): Campbell, Ian Baxter; Lambeth, Paul Francis; Naylor, Alan; Pegg, Neil Anthony  
 PATENT ASSIGNEE(S): Glaxo Group Limited, UK  
 SOURCE: PCT Int. Appl., 40 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000052008	A1	20000908	WO 1999-EP10263	19991222
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
PRIORITY APPLN. INFO.:			GB 1999-4506	A 19990227
			GB 1999-20904	A 19990903
OTHER SOURCE(S): GI		MARPAT 133:222726		



AB The title compds. [I; R0, R1 = H, halo, alkyl, etc.; R2 = halo, CN, CONR4R5, etc.; R3 = alkyl, NH2; R4, R5 = H, alkyl, (un)substituted Ph; NR4R5 = satd. 4-8 membered ring] which are potent and selective inhibitors

of COX-2 and are of use in the treatment of the pain, fever, inflammation of a variety of conditions and diseases, were prep'd. and formulated. E.g., a multi-step synthesis of I [R0 = 4-F; R1 = H; R2 = 6-CN; R3 = NH2] which showed IC50 of 21 nM against COX-2 vs. IC50 of 20,950 nM against COX-1, was given.

IT **291743-84-5P**

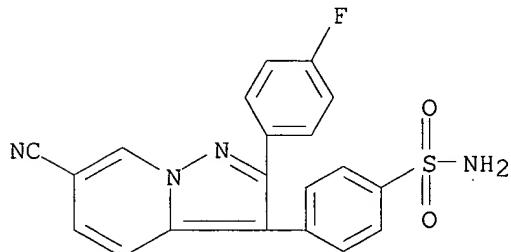
RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of pyrazolopyridines as selective inhibitors of COX-2)

RN 291743-84-5 CA

CN Benzenesulfonamide,

4-[6-cyano-2-(4-fluorophenyl)pyrazolo[1,5-a]pyridin-3-yl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT:

2

REFERENCE(S):

(1) Gauthier; US 5552422 A 1996 CA  
 (2) Glaxo Group Ltd; WO 9631509 A 1996 CA

L4 ANSWER 5 OF 5 CA

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ACCESSION NUMBER:

132:321858 CA

TITLE:

Preparation of pyrazolopyridines as selective COX-2 inhibitors

INVENTOR(S):

Campbell, Ian Baxter; Naylor, Alan

PATENT ASSIGNEE(S):

Glaxo Group Limited, UK

SOURCE:

PCT Int. Appl., 46 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

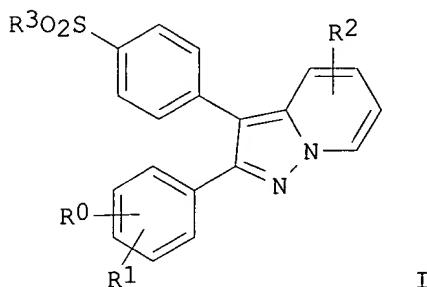
LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000026216	A1	20000511	WO 1999-EP8186	19991101
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BR 9915011	A	20010807	BR 1999-15011	19991101
EP 1127058	A1	20010829	EP 1999-955897	19991101
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
NO 2001002156	A	20010702	NO 2001-2156	20010502
PRIORITY APPLN. INFO.:			GB 1998-24062	A 19981103
			GB 1999-20909	A 19990903
			WO 1999-EP8186	W 19991101

OTHER SOURCE(S): MARPAT 132:321858  
GI

AB The title compds. [I; R0, R1 = H, halo, alkyl, etc.; R2 = H, alkyl, alkyl substituted by one or more fluorine atoms, etc.; R3 = alkyl, NH2] which are potent and selective inhibitors of COX-2 and are of use in the treatment of the pain, fever, inflammation of a variety of conditions and diseases, were prep'd. and formulated. E.g., a multi-step synthesis of I [R0 = 3-F; R1 = H; R2 = 6-CF3; R3 = NH2] which showed IC50 of 34 nM against COX-2, was given.

IT 267235-24-5P

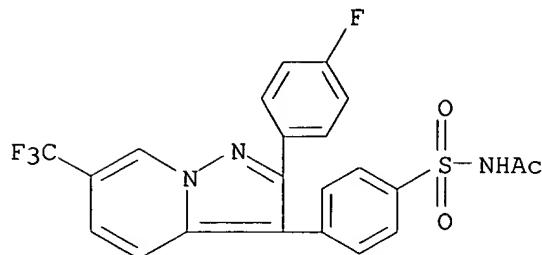
RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of pyrazolopyridines as selective COX-2 inhibitors)

RN 267235-24-5 CA

CN Acetamide, N-[[4-[2-(4-fluorophenyl)-6-(trifluoromethyl)pyrazolo[1,5-

a]pyridin-3-yl]phenyl]sulfonyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT:

8

REFERENCE(S):

- (1) Carter, J; EXPERT OPINION ON THERAPEUTIC PATENTS 1998, V8(1), P21 CA
- (2) Glaxo Group Ltd; WO 9631509 A 1996 CA
- (3) Merck Frosst Canada Inc; WO 9606840 A 1996 CA
- (4) Merck Frosst Canada Inc; WO 9621667 A 1996 CA
- (5) Naylor, A; WO 9912930 A 1999 CA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

=&gt; file caold

COST IN U.S. DOLLARS

	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	23.84	157.86

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-2.80	-2.80

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FILE LAST UPDATED: 01 May 1997 (19970501/UP)

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This file supports REGISTRY for direct browsing and searching of all substance data from the REGISTRY file. Enter HELP FIRST for more information.

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FILE 'REGISTRY' ENTERED AT 15:07:30 ON 20 NOV 2001  
L1 STRUCTURE uploaded  
L2 8 S L1  
L3 160 S L1 FULL

FILE 'CA' ENTERED AT 15:08:27 ON 20 NOV 2001  
L4 5 S L3  
L5 0 S L4 AND BAXTER, I?/AU  
L6 0 S L4 AND PD < DECEMBER 1998

FILE 'CAOLD' ENTERED AT 15:10:02 ON 20 NOV 2001

=> s 13

L7 0 L3

=>

=>  
Executing the logoff script...

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-2.80

SESSION WILL BE HELD FOR 60 MINUTES  
STN INTERNATIONAL SESSION SUSPENDED AT 15:10:22 ON 20 NOV 2001

---Logging off of STN---

END

Unable to generate the STN prompt.  
Exiting the script...